

FILE 'REGISTRY' ENTERED AT 10:27:10 ON 26 APR 2007  
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STRUCTURE FILE UPDATES: 25 APR 2007 HIGHEST RN 932710-95-7  
DICTIONARY FILE UPDATES: 25 APR 2007 HIGHEST RN 932710-95-7

New CAS Information Use Policies, enter HELP USAGETERMS for details.

TSCA INFORMATION NOW CURRENT THROUGH December 2, 2006

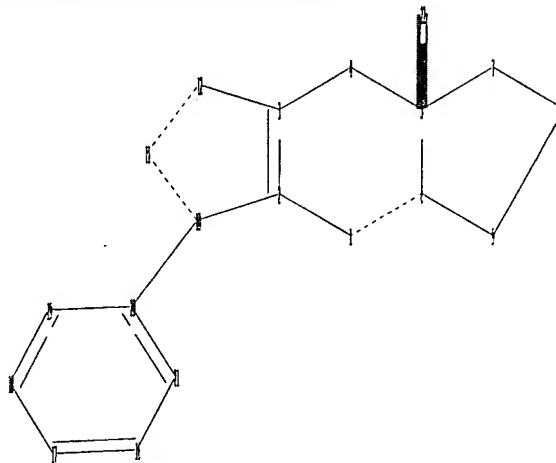
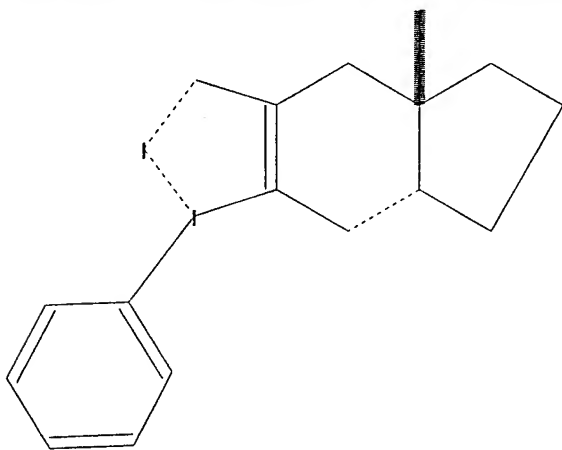
Please note that search-term pricing does apply when  
conducting SmartSELECT searches.

REGISTRY includes numerically searchable data for experimental and  
predicted properties as well as tags indicating availability of  
experimental property data in the original document. For information  
on property searching in REGISTRY, refer to:

<http://www.cas.org/support/stngen/stndoc/properties.html>

=>

Uploading C:\Program Files\Stnexp\Queries\10.544899\electd group.str



chain nodes :

22

ring nodes :

1 2 3 4 5 6 7 8 9 10 11 12 16 17 18 19 20 21

chain bonds :

5-22 10-20

ring bonds :

1-2 1-6 2-3 2-10 3-4 3-12 4-5 5-6 5-7 6-9 7-8 8-9 10-11 11-12 16-17 16-21 17-18  
18-19 19-20 20-21

exact/norm bonds :

1-6 8-9 10-11 10-20 11-12

exact bonds :

1-2 2-3 2-10 3-4 3-12 4-5 5-6 5-7 5-22 6-9 7-8

normalized bonds :

16-17 16-21 17-18 18-19 19-20 20-21

isolated ring systems :

containing 1 :

G1:C,N

Match level :

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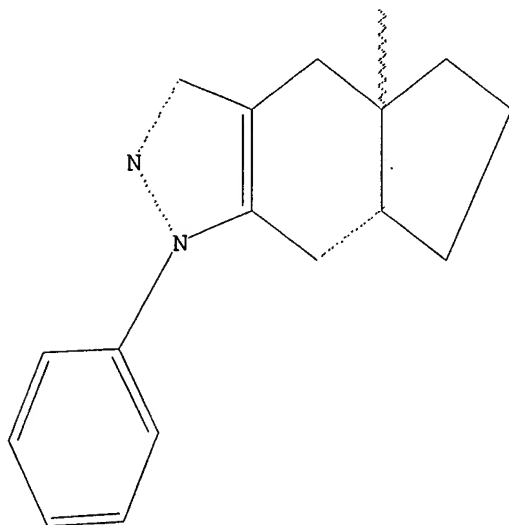
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12:Atom 16:Atom 17:Atom 18:Atom 19:Atom 20:Atom 21:Atom 22:CLASS

L1 STRUCTURE UPLOADED

=> d

L1 HAS NO ANSWERS

L1 STR



G1 C,N

Structure attributes must be viewed using STN Express query preparation.

=> s l1 sss sam

SAMPLE SEARCH INITIATED 10:28:00 FILE 'REGISTRY'

SAMPLE SCREEN SEARCH COMPLETED - 110 TO ITERATE

100.0% PROCESSED 110 ITERATIONS

14 ANSWERS

SEARCH TIME: 00.00.01

FULL FILE PROJECTIONS: ONLINE \*\*COMPLETE\*\*  
BATCH \*\*COMPLETE\*\*

PROJECTED ITERATIONS: 1571 TO 2829

PROJECTED ANSWERS: 56 TO 504

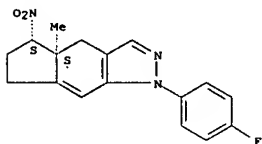
L2 14 SEA SSS SAM L1

=> d scan

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L2 14 ANSWERS REGISTRY COPYRIGHT 2007 ACS on STN  
IN Cyclopent[*f*]indazole, 1-(4-fluorophenyl)-1,4,4a,5,6,7-hexahydro-4a-methyl-  
5-nitro-, (4a*S*,5*S*)- (9CI)  
MF C17 H16 F N3 O2

Absolute stereochemistry.

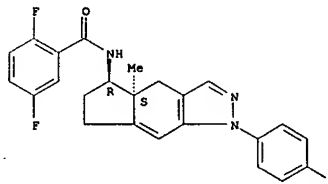


\*\*PROPERTY DATA AVAILABLE IN THE 'PROP' FORMAT\*\*

HOW MANY MORE ANSWERS DO YOU WISH TO SCAN? (1):1

L2 14 ANSWERS REGISTRY COPYRIGHT 2007 ACS on STN  
IN Benzamide, 2,5-difluoro-N-[(4*aS*,5*R*)-1-(4-fluorophenyl)-1,4,4a,5,6,7-  
hexahydro-4a-methylcyclopent[*f*]indazol-5-yl]- (9CI)  
MF C24 H20 F3 N3 O

Absolute stereochemistry.

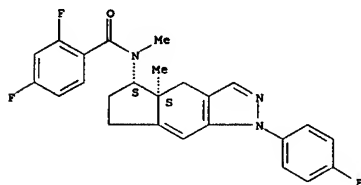


\*\*PROPERTY DATA AVAILABLE IN THE 'PROP' FORMAT\*\*

HOW MANY MORE ANSWERS DO YOU WISH TO SCAN? (1):1

L2 14 ANSWERS REGISTRY COPYRIGHT 2007 ACS on STN  
IN Benzamide, 2,4-difluoro-N-[(4*aS*,5*S*)-1-(4-fluorophenyl)-1,4,4a,5,6,7-  
hexahydro-4a-methylcyclopent[*f*]indazol-5-yl]-N-methyl- (9CI)  
MF C25 H22 F3 N3 O

Absolute stereochemistry.



\*\*PROPERTY DATA AVAILABLE IN THE 'PROP' FORMAT\*\*

HOW MANY MORE ANSWERS DO YOU WISH TO SCAN? (1):0

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=> file hcaplus

COST IN U.S. DOLLARS

SINCE FILE

TOTAL

ENTRY

SESSION

FULL ESTIMATED COST

0.90

1.11

FILE 'HCAPLUS' ENTERED AT 10:28:33 ON 26 APR 2007

USE IS SUBJECT TO THE TERMS OF YOUR STN CUSTOMER AGREEMENT.

PLEASE SEE "HELP USAGETERMS" FOR DETAILS.

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FILE COVERS 1907 - 26 Apr 2007 VOL 146 ISS 18

FILE LAST UPDATED: 25 Apr 2007 (20070425/ED)

New CAS Information Use Policies, enter HELP USAGETERMS for details.

This file contains CAS Registry Numbers for easy and accurate substance identification.

=> s l2

L3 3 L2

=> d ibib abs hitstr tot

L3 ANSWER 1 OF 3 HCAPLUS COPYRIGHT 2007 ACS on STN (Continued)  
 PREP. of pyrazoles bearing annulated spirocyclic hydronaphthalene  
 derivs. as glucocorticoid receptor modulators)  
 RN 786706-88-5 HCAPLUS  
 CN Cyclopent[*f*]indazole, 1-(4-fluorophenyl)-1,4,4a,5,6,7-hexahydro-4a-methyl-  
 5-nitro-, (4a*S*,5*S*)- (9CI) (CA INDEX NAME)

INVENTOR(S): Ali, Amjad; Balkovec, James M.; Beresin, Richard;  
 Colletti, Steven L.; Graham, Donald W.; Patel, Gool  
 F.; Smith, Cameron J.  
 PATENT ASSIGNEE(S): Merck & Co., Inc., USA  
 SOURCE: PCT Int. Appl., 201 pp.  
 CODEN: PIXXD2  
 DOCUMENT TYPE: Patent  
 LANGUAGE: English  
 FAMILY ACC. NUM. COUNT: 1  
 PATENT INFORMATION:

PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
WO 2004093805	A2	20041104	WO 2004-US12102	20040419
WO 2004093805	A3	20051208		
W:	AE, AG, AL, AM, AT, AU, AZ, BA, BB, BG, BR, BW, BY, BZ, CA, CH, CN, CO, CR, CU, CZ, DE, DK, DM, DZ, EC, EE, EG, ES, FI, GB, GD, GE, GH, GM, GR, GU, ID, IL, IN, IS, JP, KE, KG, KP, KR, KZ, LC, LK, LR, LS, LT, LU, LV, MA, MD, MG, MK, MN, MW, MX, MZ, NA, NI, NO, NZ, OM, PG, PH, PL, PT, RO, RU, SC, SD, SE, SG, SK, SL, SY, TJ, TM, TN, TR, TT, UA, UG, US, VC, VN, YU, ZA, ZM, ZW			
RW:	BW, GH, GM, KE, LS, MW, MZ, SD, SI, SZ, TG, UG, ZM, ZW, AM, AZ, BY, KG, KZ, MD, RU, TJ, TM, AT, BE, BG, CH, CY, CZ, DE, DK, EE, ES, FI, FR, GB, GR, HU, IE, IT, LU, MC, NL, PL, PT, RO, SE, SI, SK, TR, BF, BJ, CF, CG, CI, CM, GA, GN, GQ, GW, ML, MR, NE, SN, TD, TG			
AU 2004232301	A1	20041104	AU 2004-232301	20040419
CA 2522946	A1	20041104	CA 2004-2522946	20040419
EP 1617806	A2	20060125	EP 2004-760029	20040419
R:	AT, BE, CH, DE, DK, ES, FR, GB, GR, IT, LI, LU, NL, SE, MC, PT, IE, SI, LT, LV, FI, RO, MK, CY, AL, TR, BG, CZ, EE, HU, PL, SK, HR			
CN 1809347	A	20060726	CN 2004-80017051	20040419
JP 2006524251	T	20061026	JP 2006-513140	20040419
US 2006217563	A1	20060928	US 2005-551933	20051004
PRIORITY APPL. INFO.:			US 2003-464784P	P 20030423
			WO 2004-US12102	W 20040419

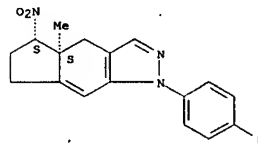
OTHER SOURCE(S): MARPAT 141:395547  
 GI

\* STRUCTURE DIAGRAM TOO LARGE FOR DISPLAY - AVAILABLE VIA OFFLINE PRINT \*

AB Title compds. I [Ring A = carbocyclic or heterocyclic; m = 0-3; n = 0-2; R1 = (un)substituted-alkyl, -alkenyl, -alkynyl, -cycloalkyl, etc.; R2 and R3 independently = H, halo, alkyl, aryl, etc.; R4 = OH, CO2H, (un)substituted-alkyl, -Ph, etc.], as well as their pharmaceutically acceptable salts or hydrates thereof, are prepared and disclosed as selective glucocorticoid receptor ligands for treating a variety of autoimmune and inflammatory diseases or conditions. Thus, e.g., II was prepared via spirocyclization of III (preparation given) with Et  $\alpha$ -bromomethyl acrylate. In human glucocorticoid receptor assays, I demonstrated a range of GR affinity with IC50 values between 10  $\mu$ M and 1 nM. Pharmaceutical compds. and methods of use are also included.

IT 786706-88-5P 786706-89-6P  
 RL: RCT (Reactant); SPN (Synthetic preparation); PREP (Preparation); RACT (Reactant or reagent)

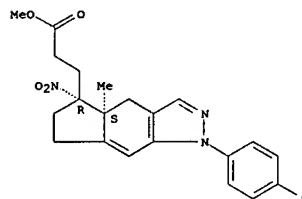
Absolute stereochemistry.



RN 786706-89-6 HCAPLUS

CN Cyclopent[*f*]indazole-5-propanoic acid, 1-(4-fluorophenyl)-1,4,4a,5,6,7-hexahydro-4a-methyl-5-nitro-, methyl ester, (4a*S*,5*R*)- (9CI) (CA INDEX NAME)

Absolute stereochemistry.



L3 ANSWER 2 OF 3 HCAPLUS COPYRIGHT 2007 ACS on STN (Continued)  
 PREP. of pyrazoles bearing annulated spirocyclic hydronaphthalene  
 derivs. as glucocorticoid receptor modulators)  
 RN 786706-88-5 HCAPLUS  
 CN Cyclopent[*f*]indazole, 1-(4-fluorophenyl)-1,4,4a,5,6,7-hexahydro-4a-methyl-  
 5-nitro-, (4a*S*,5*S*)- (9CI) (CA INDEX NAME)

INVENTOR(S): Ali, Amjad; Beresin, Richard; Colletti, Steven L.;  
 Graham, Donald W.; Tata, James R.; Thompson,  
 Christopher F.  
 PATENT ASSIGNEE(S): Merck & Co. Inc., USA  
 SOURCE: PCT Int. Appl., 105 pp.  
 CODEN: PIXXD2  
 DOCUMENT TYPE: Patent  
 LANGUAGE: English  
 FAMILY ACC. NUM. COUNT: 1  
 PATENT INFORMATION:

PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
WO 2004075840	A2	20040910	WO 2004-US5199	20040220
WO 2004075840	A3	20050203		
WO 2004075840	A9	20050804		
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RW:	BW, GH, GM, KE, LS, MW, MZ, SD, SI, SZ, TG, UG, ZM, ZW, AM, AZ, BY, KG, KZ, MD, RU, TJ, TM, AT, BE, BG, CH, CY, CZ, DE, DK, EE, ES, FI, FR, GB, GR, HU, IE, IT, LU, MC, NL, PT, RO, SE, SI, SK, TR, BF, BJ, CF, CG, CI, CM, GA, GN, GQ, GW, ML, MR, NE, SN, TD, TG			
AU 2004216182	A1	20040910	AU 2004-216182	20040220
CA 2516684	A1	20040910	CA 2004-2516684	20040220
EP 1599201	A2	20051130	EP 2004-713398	20040220
R:	AT, BE, CH, DE, DK, ES, FR, GB, GR, IT, LI, LU, NL, SE, MC, PT, IE, SI, LT, LV, FI, RO, MK, CY, AL, TR, BG, CZ, EE, HU, SK			
JP 2006518752	T	20060817	JP 2006-503780	20040220
US 2006074120	A1	20060406	US 2005-544899	20050808
PRIORITY APPL. INFO.:			US 2003-450811P	P 20030225
			WO 2004-US5199	W 20040220

OTHER SOURCE(S): MARPAT 141:260743  
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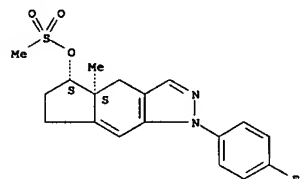
L3 ANSWER 2 OF 3 HCAPLUS COPYRIGHT 2007 ACS on STN (Continued)

AB Title compds. represented by the formula I [wherein J = NR1, CR1R2; K = NR3, CR3R4; L = NR5, CR5R6; X = hydroxy, alkoxy, carbamoyl, etc.; R1-R6 = independently H, halo, (cyclo)alkyl, etc.; R7 = H, hydroxy, alkoxy, aryl, etc.; R8 = (cyclo)alkyl, alkenyl, alkynyl, etc.; R9, R10 = independently halo, hydroxy, alkyl, alkenyl, alkoxy; n = 0-2; and pharmaceutically acceptable salts or hydrates thereof] were prepared as selective non-steroidal glucocorticoid receptor modulators. For example, II was given in a multi-steps synthesis starting from 1-(4-fluorophenyl)-4,4a,6,7-tetrahydro-4a-methyl-cyclopent[*f*]indazol-5(1H)-one reacting with phenylethynylmagnesium bromide. I showed affinity of glucocorticoid receptor with IC50 values between 10  $\mu$ M and 1 nM. Thus, I and their pharmaceutical compds. are useful for the treatment of a variety of autoimmune and inflammatory diseases or conditions.

IT 754237-88-2P  
 RL: PAC (Pharmacological activity); RCT (Reactant); SPN (Synthetic preparation); THU (Therapeutic use); BIOL (Biological study); PREP (Preparation); RACT (Reactant or reagent); USES (Uses)  
 (preparation of cyclopent[*f*]indazol-5-yl and benz[*f*]indazol-5-yl derivs. as selective non-steroidal glucocorticoid receptor modulators)

RN 754237-88-2 HCAPLUS  
 CN Cyclopent[*f*]indazol-5-ol, 1-(4-fluorophenyl)-1,4,4a,5,6,7-hexahydro-4a-methyl-, methanesulfonate (ester), (4a*S*,5*S*)- (9CI) (CA INDEX NAME)

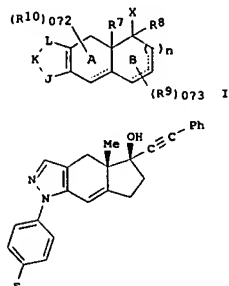
Absolute stereochemistry.

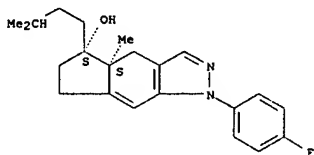


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 754238-00-1P 754238-30-7P 754238-35-2P  
 RL: PAC (Pharmacological activity); SPN (Synthetic preparation); THU (Therapeutic use); BIOL (Biological study); PREP (Preparation); USES (Uses)  
 (preparation of cyclopent[*f*]indazol-5-yl and benz[*f*]indazol-5-yl derivs. as selective non-steroidal glucocorticoid receptor modulators)

RN 754236-00-5 HCAPLUS  
 CN Cyclopent[*f*]indazol-5-ol, 1-(4-fluorophenyl)-1,4,4a,5,6,7-hexahydro-4a-methyl-5-(3-methylbutyl)-, (4a*S*,5*S*)- (9CI) (CA INDEX NAME)

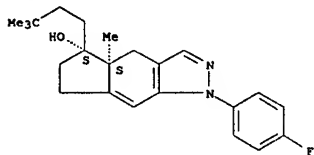
Absolute stereochemistry.





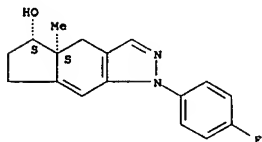
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CN Cyclopent[*f*]indazol-5-yl, 5-[(3,3-dimethylbutyl)-1-(4-fluorophenyl)-1,4,4a,5,6,7-hexahydro-4a-methyl-, (4a*S*,5*S*)- (9*CI*) (CA INDEX NAME)

Absolute stereochemistry.



RN 754236-65-2 HCAPLUS  
CN Cyclopent[*f*]indazol-5-yl, 1-(4-fluorophenyl)-1,4,4a,5,6,7-hexahydro-4a-methyl-, (4a*S*,5*S*)- (9*CI*) (CA INDEX NAME)

Absolute stereochemistry.

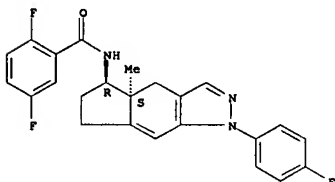


RN 754236-75-4 HCAPLUS  
CN Cyclopent[*f*]indazole, 5-[(2-(3-chlorophenyl)ethyl)-1-(4-fluorophenyl)-1,4,4a,5,6,7-hexahydro-4a-methyl-5-propoxy-, (4a*S*,5*S*)- (9*CI*) (CA INDEX NAME)

Absolute stereochemistry.

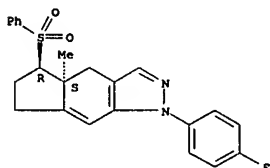
RN 754238-00-1 HCAPLUS  
CN Benzamide, 2,5-difluoro-N-[(4a*S*,5*R*)-1-(4-fluorophenyl)-1,4,4a,5,6,7-hexahydro-4a-methylcyclopent[*f*]indazol-5-yl]- (9*CI*) (CA INDEX NAME)

Absolute stereochemistry.



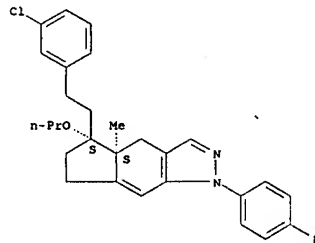
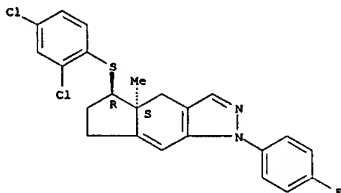
RN 754238-30-7 HCAPLUS  
CN Cyclopent[*f*]indazole, 1-(4-fluorophenyl)-1,4,4a,5,6,7-hexahydro-4a-methyl-5-(phenylsulfonyl)-, (4a*S*,5*R*)- (9*CI*) (CA INDEX NAME)

Absolute stereochemistry.



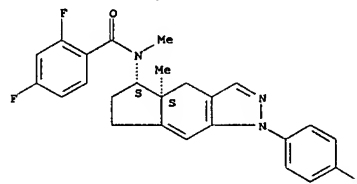
RN 754238-35-2 HCAPLUS  
CN Cyclopent[*f*]indazole, 5-[(2,4-dichlorophenylthio)-1-(4-fluorophenyl)-1,4,4a,5,6,7-hexahydro-4a-methyl-, (4a*S*,5*R*)- (9*CI*) (CA INDEX NAME)

Absolute stereochemistry.



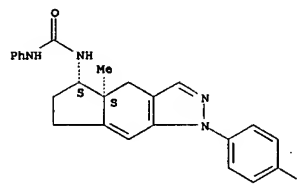
RN 754237-30-4 HCAPLUS  
CN Benzamide, 2,4-di-fluoro-N-[(4a*S*,5*S*)-1-(4-fluorophenyl)-1,4,4a,5,6,7-hexahydro-4a-methylcyclopent[*f*]indazol-5-yl]-N-methyl- (9*CI*) (CA INDEX NAME)

Absolute stereochemistry.



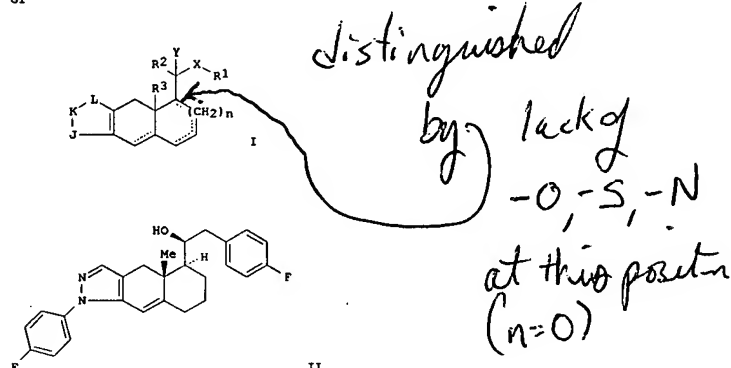
RN 754237-65-5 HCAPLUS  
CN Urea, N-[(4a*S*,5*S*)-1-(4-fluorophenyl)-1,4,4a,5,6,7-hexahydro-4a-methylcyclopent[*f*]indazol-5-yl]-N'-phenyl- (9*CI*) (CA INDEX NAME)

Absolute stereochemistry.



ACCESSION NUMBER: 2003:836773 HCAPLUS  
DOCUMENT NUMBER: 139:323524  
TITLE: Preparation of 1H-Benzo[*f*]indazol-5-yl derivatives as selective glucocorticoid receptor modulators  
INVENTOR(S): Ali, Amjad; Balkovec, James M.; Graham, Donald W.; Thompson, Christopher F.; Quraishi, Nazia  
PATENT ASSIGNEE(S): Merck & Co., Inc., USA  
SOURCE: PCT Int. Appl., 233 pp.  
DOCUMENT TYPE: CODEN: PIXXD2  
LANGUAGE: Patent  
FAMILY ACC. NUM. COUNT: English  
PATENT INFORMATION: 1

PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
WO 2003086294	A2	20031023	WO 2003-US10867	20030408
WO 2003086294	A3	20040715		
W: AE, AG, AL, AM, AT, AU, AZ, BA, BB, BG, BR, BY, BZ, CA, CH, CN, CO, CR, CU, CZ, DE, DK, DM, DZ, EC, EE, ES, FI, GB, GD, GE, GH, GM, HR, HU, ID, IL, IN, IS, JP, KE, KG, KR, KZ, LC, LK, LR, LS, LT, LU, LV, MA, MD, MG, MK, MN, MW, MX, MY, NZ, OM, PH, PL, PT, RO, RU, SC, SD, SE, SG, SK, SL, TJ, TM, TN, TR, TT, TZ, UA, UG, US, UZ, VC, VN, YU, ZA, ZM, ZW				
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CA 2481320	A1	20031023	CA 2003-2481320	20030408
AU 2003221706	A1	20031027	AU 2003-221706	20030408
EP 1496892	A2	20050119	EP 2003-718285	20030408
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JP 2005528385	T	20050922	JP 2003-583321	20030408
US 2005256315	A1	20051117	US 2005-508897	20050428
PRIORITY APPLN. INFO.: US 2002-371948P P 20020411				
OTHER SOURCE(S): MARPAT 139:323524				
GI				

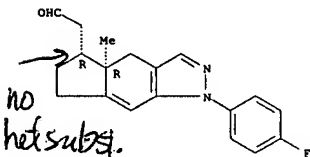


AB Benzindazoles I (n = 0-2; J, K, L = (un)substituted CH<sub>2</sub>, NH; X = bond, CO, (un)substituted NH, NHCO, 1,1-cyclopropanediyl; R<sub>1</sub>, R<sub>2</sub> = H, (un)substituted alkyl, alkenyl, alkynyl, cycloalkyl, alkoxy, alkylthio, alkylsulfinyl, alkylsulfonyl, aryl, aralkyl, heterocyclic, aryloxy, aryloxy, OH; R<sub>3</sub> = H, (un)substituted OH, alkyl, aryl, aralkyl; Y = H, (un)substituted OH, SH, S(O)H, SO<sub>2</sub>H, CH<sub>2</sub>, NH<sub>2</sub>, SO<sub>2</sub>NH<sub>2</sub>, CO<sub>2</sub>H, NO<sub>2</sub>, acyl, CN, halogen; and the carbocyclic rings may be further substituted) were prepared for use as selective glucocorticoid receptor ligands for treating a variety of autoimmune and inflammatory diseases or conditions (no data). Thus, Wieland-Miescher ketone was ketalized, hydroxymethylenated, cyclized with 4-FC6H<sub>4</sub>NNNH<sub>2</sub>, deketalized, treated with Ph<sub>3</sub>P+CH<sub>2</sub>OMe Cl<sup>-</sup>, and subjected to Grignard reaction with 4-FC6H<sub>4</sub>MgCl to give the benzindazole II.

IT 614763-27-8P  
RL: RCT (Reactant); SPN (Synthetic preparation); PREP (Preparation); RACT (Reactant or reagent)  
(preparation of 1H-benzo[f]indazol-5-yl derivs. as selective glucocorticoid receptor modulators)

RN 614763-27-8 HCAPLUS  
CN Cyclopent[f]indazole-5-acetaldehyde, 1-(4-fluorophenyl)-1,4,4a,5,6,7-hexahydro-4a-methyl-, (4aR,5R)- (9CI) (CA INDEX NAME)

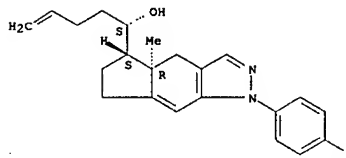
Absolute stereochemistry.



IT 614761-89-6P  
RL: SPN (Synthetic preparation); THU (Therapeutic use); BIOL (Biological study); PREP (Preparation); USES (Uses)  
(preparation of 1H-benzo[f]indazol-5-yl derivs. as selective glucocorticoid receptor modulators)

RN 614761-89-6 HCAPLUS  
CN Cyclopent[f]indazole-5-methanol, α-3-butenyl-1-(4-fluorophenyl)-1,4,4a,5,6,7-hexahydro-4a-methyl-, (αS,4aR,5S)- (9CI) (CA INDEX NAME)

Absolute stereochemistry.



=&gt; file reg

COST IN U.S. DOLLARS	SINCE FILE ENTRY	TOTAL SESSION
FULL ESTIMATED COST	34.01	35.12
DISCOUNT AMOUNTS (FOR QUALIFYING ACCOUNTS)	SINCE FILE ENTRY	TOTAL SESSION
CA SUBSCRIBER PRICE	-2.34	-2.34

FILE 'REGISTRY' ENTERED AT 10:32:56 ON 26 APR 2007  
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STRUCTURE FILE UPDATES: 25 APR 2007 HIGHEST RN 932710-95-7  
 DICTIONARY FILE UPDATES: 25 APR 2007 HIGHEST RN 932710-95-7

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<http://www.cas.org/support/stngen/stndoc/properties.html>

=&gt; s ll sss full

FULL SEARCH INITIATED 10:33:05 FILE 'REGISTRY'  
 FULL SCREEN SEARCH COMPLETED - 2196 TO ITERATE

100.0% PROCESSED 2196 ITERATIONS 272 ANSWERS  
 SEARCH TIME: 00.00.01

L4 272 SEA SSS FUL L1

=&gt; file hcaplus

COST IN U.S. DOLLARS	SINCE FILE ENTRY	TOTAL SESSION
FULL ESTIMATED COST	172.10	207.22
DISCOUNT AMOUNTS (FOR QUALIFYING ACCOUNTS)	SINCE FILE ENTRY	TOTAL SESSION
CA SUBSCRIBER PRICE	0.00	-2.34

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FILE COVERS 1907 - 26 Apr 2007 VOL 146 ISS 18  
FILE LAST UPDATED: 25 Apr 2007 (20070425/ED)

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This file contains CAS Registry Numbers for easy and accurate  
substance identification.

=> s l4

L5 4 L4

=> s l5 not l3

L6 1 L5 NOT L3

=> d ibib abs hitstr

L6 ANSWER 1 OF 1 HCAPLUS COPYRIGHT 2007 ACS on STN

ACCESSION NUMBER: 2005:461153 HCAPLUS

DOCUMENT NUMBER: 143:125827

TITLE: Novel ketal ligands for the glucocorticoid receptor:

in vitro and in vivo activity

AUTHOR(S): Smith, Cameron J.; Ali, Amjad; Balkovec, James M.;  
 Graham, Donald W.; Hammond, Milton L.; Patel, Gool F.;  
 Rouen, Gregory P.; Smith, Scott K.; Tata, James R.;  
 Einstein, Monica; Ge, Lan; Harris, Georgianna S.;  
 Kelly, Theresa M.; Mazur, Paul; Thompson, Chris M.;  
 Wang, Chuanlin F.; Williamson, Joanne M.; Miller,  
 Douglas K.; Pandit, Shilpa; Santoro, Joseph C.;  
 Sitlani, Ayesha; Yamin, Ting-ting D.; O'Neill, Edward  
 A.; Zaller, Dennis M.; Carballo-Jane, Ester; Forrest,  
 Michael J.; Luell, Silvi

CORPORATE SOURCE: Department of Medicinal Chemistry, Merck Research

Laboratories, Rahway, NJ, 07065, USA

SOURCE: Bioorganic &amp; Medicinal Chemistry Letters (2005),

15(11), 2926-2931

CODEN: BMCL88; ISSN: 0960-894X

PUBLISHER: Elsevier B.V.

DOCUMENT TYPE: Journal

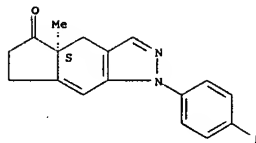
LANGUAGE: English

OTHER SOURCE(S): CASREACT 143:125827

GI

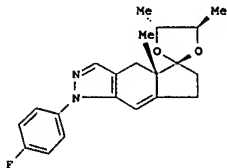
L6 ANSWER 1 OF 1 HCAPLUS COPYRIGHT 2007 ACS on STN

(Continued)



REFERENCE COUNT:

36

THERE ARE 36 CITED REFERENCES AVAILABLE FOR THIS  
RECORD. ALL CITATIONS AVAILABLE IN THE RE FORMAT

I

AB A novel series of selective ligands for the human glucocorticoid receptor is described. Structure-activity studies focused on variation of B-ring size, ketal ring size, and ketal substitution. These analogs were found to be potent and selective ligands for GR and have partial agonist profiles in functional assays for transactivation (TAT, GS) and transrepression (IL-6). Of these compds., three were evaluated further in a mouse LPS-induced TNF- $\alpha$  secretion model. Compound (I) had an ED50 of 14.1 mg/kg compared with 0.5 mg/kg for prednisolone in the same assay.

IT 614763-02-9P

RL: RCT (Reactant); SPN (Synthetic preparation); PREP (Preparation); RACT

(Reactant or reagent)

(novel ketal ligands for glucocorticoid receptor and in vitro and in

vivo activity)

RN 614763-02-9 HCAPLUS

CN Cyclopent[*f*]indazol-5(1H)-one, 1-(4-fluorophenyl)-4,4a,6,7-tetrahydro-4a-methyl-, (4aS)- (9CI) (CA INDEX NAME)

Absolute stereochemistry.

Robert Havlin - 10.544899

=> log hold

COST IN U.S. DOLLARS

SINCE FILE

TOTAL

ENTRY

SESSION

FULL ESTIMATED COST

10.47

217.69

DISCOUNT AMOUNTS (FOR QUALIFYING ACCOUNTS)

SINCE FILE

TOTAL

ENTRY

SESSION

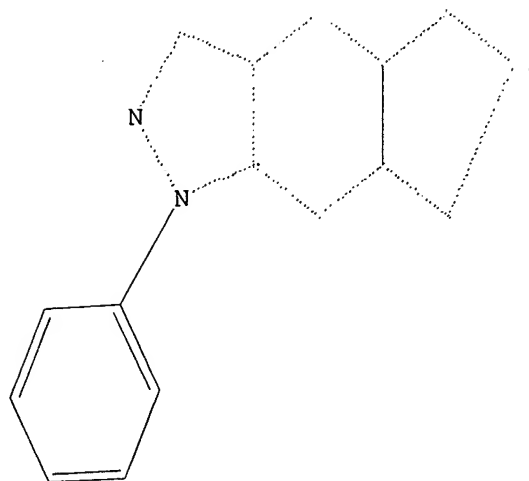
CA SUBSCRIBER PRICE

-0.78

-3.12

SESSION WILL BE HELD FOR 120 MINUTES

STN INTERNATIONAL SESSION SUSPENDED AT 10:34:35 ON 26 APR 2007



G1 C,N

G2 O,S,N

Structure attributes must be viewed using STN Express query preparation.

=> s l11 sss full

FULL SEARCH INITIATED 10:47:54 FILE 'REGISTRY'

FULL SCREEN SEARCH COMPLETED - 11557 TO ITERATE

100.0% PROCESSED 11557 ITERATIONS

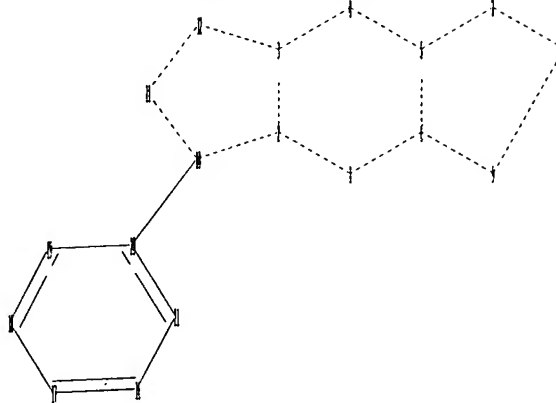
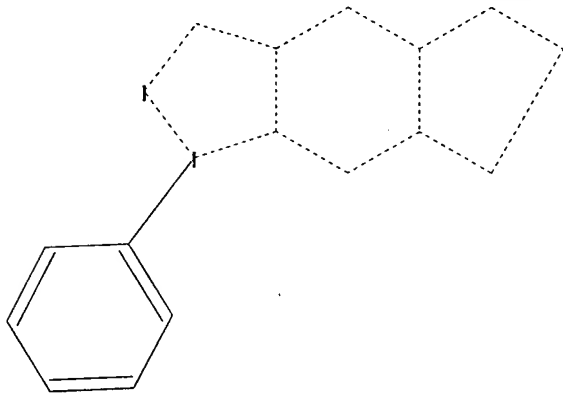
273 ANSWERS

SEARCH TIME: 00.00.01

L12 273 SEA SSS FUL L11

=>

Uploading C:\Program Files\Stnexp\Queries\10.544899\electd group D.str



ring nodes :

1 2 3 4 5 6 7 8 9 10 11 12 16 17 18 19 20 21

chain bonds :

10-20

ring bonds :

1-2 1-6 2-3 2-10 3-4 3-12 4-5 5-6 5-7 6-9 7-8 8-9 10-11 11-12 16-17 16-21 17-18  
18-19 19-20 20-21

exact/norm bonds :

1-2 1-6 2-3 2-10 3-4 3-12 4-5 5-6 5-7 6-9 7-8 8-9 10-11 10-20 11-12

normalized bonds :

16-17 16-21 17-18 18-19 19-20 20-21

isolated ring systems :

containing 1 :

G1:C,N

G2:O,S,N

Match level :

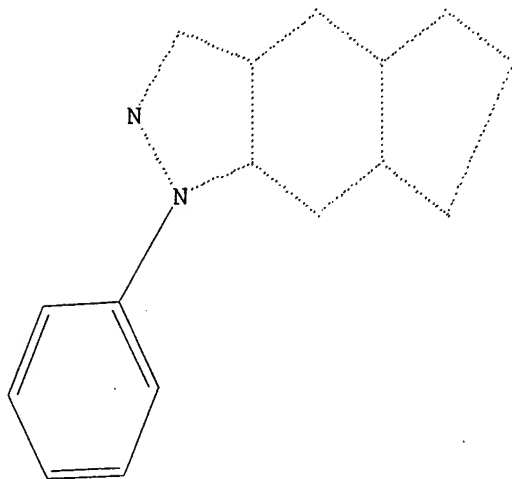
1:Atom 2:Atom 3:Atom 4:Atom 5:Atom 6:Atom 7:Atom 8:Atom 9:Atom 10:Atom 11:Atom  
12:Atom 16:Atom 17:Atom 18:Atom 19:Atom 20:Atom 21:Atom

L13 STRUCTURE UPLOADED

=> d

L13 HAS NO ANSWERS

L13 STR



G1 C,N

G2 O,S,N

Structure attributes must be viewed using STN Express query preparation.

=> s l13 sss sam

SAMPLE SEARCH INITIATED 10:49:03 FILE 'REGISTRY'

SAMPLE SCREEN SEARCH COMPLETED - 601 TO ITERATE

100.0% PROCESSED 601 ITERATIONS

14 ANSWERS

SEARCH TIME: 00.00.01

FULL FILE PROJECTIONS: ONLINE \*\*COMPLETE\*\*

BATCH \*\*COMPLETE\*\*

PROJECTED ITERATIONS: 10550 TO 13490

PROJECTED ANSWERS: 56 TO 504

L14 14 SEA SSS SAM L13

=> s l13 sss full

FULL SEARCH INITIATED 10:49:08 FILE 'REGISTRY'

FULL SCREEN SEARCH COMPLETED - 11557 TO ITERATE

100.0% PROCESSED 11557 ITERATIONS

277 ANSWERS

SEARCH TIME: 00.00.01

L15 277 SEA SSS FUL L13

=> file hcaplus

Robert Havlin - 10.544899

COST IN U.S. DOLLARS	SINCE FILE ENTRY	TOTAL SESSION
FULL ESTIMATED COST	344.65	739.64
DISCOUNT AMOUNTS (FOR QUALIFYING ACCOUNTS)	SINCE FILE ENTRY	TOTAL SESSION
CA SUBSCRIBER PRICE	0.00	-3.12

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FILE COVERS 1907 - 26 Apr 2007 VOL 146 ISS 18  
FILE LAST UPDATED: 25 Apr 2007 (20070425/ED)

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This file contains CAS Registry Numbers for easy and accurate substance identification.

=> s l15  
L16 7 L15  
  
=> s l16 not l5  
L17 3 L16 NOT L5  
  
=> d ibib abs hitstr tot

L17 ANSWER 1 OF 3 HCAPLUS COPYRIGHT 2007 ACS on STN

ACCESSION NUMBER: 1991:607969 HCAPLUS

DOCUMENT NUMBER: 115:207969

TITLE: Condensed pyridazines. VIII. Reaction of diazolopyridazines with ynamine. Formation of benzodiazoles and diazolidiazocines

AUTHOR(S): Oishi, Etsuo; Teido, Naokata; Miyashita, Akira; Sato, Susumu; Ohta, Syouji; Ishida, Hitoshi; Tanji, Kenichi; Higashino, Takeo

CORPORATE SOURCE: Sch. Pharm. Sci., Univ. Shizuoka, Shizuoka, 422, Japan

SOURCE: Chemical &amp; Pharmaceutical Bulletin (1991), 39(7), 1713-18

CODEN: CPBTAL; ISSN: 0009-2363

DOCUMENT TYPE: Journal

LANGUAGE: English

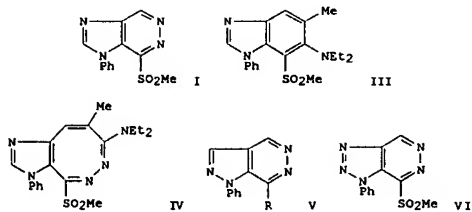
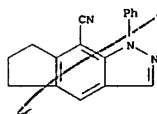
OTHER SOURCE(S): CASREACT 115:207969

GI

L17 ANSWER 1 OF 3 HCAPLUS COPYRIGHT 2007 ACS on STN

(CA INDEX NAME)

(Continued)



AB Reaction of imidazopyridazine I with MeC.tplbond.CNt2 (II) resulted in the formation of the benzimidazole III and the 3H-imidazo[4,5-d][1,2]diazocine IV through the [4+2]- and [2+2]cycloadducts. Similarly, reaction of the pyrazolopyridazines V (R = CN, SO<sub>2</sub>Me) with II gave the corresponding indazoles and pyrazolodiazocines. A similar reaction was also found to proceed between (methylsulfonyl)triazolopyridazines VI and II affording the corresponding benzotriazole and triazolodiazocine.

IT 136819-80-2P 136819-81-3P

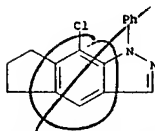
RL: SPN (Synthetic preparation); PREP (Preparation)

(preparation of)

RN 136819-80-2 HCAPLUS

CN Cyclopent[1,2]indazole, 8-chloro-1,5,6,7-tetrahydro-1-phenyl- (9CI) (CA

INDEX NAME)



RN 136819-81-3 HCAPLUS

CN Cyclopent[1,2]indazole-8-carbonitrile, 1,5,6,7-tetrahydro-1-phenyl- (9CI)

L17 ANSWER 2 OF 3 HCAPLUS COPYRIGHT 2007 ACS on STN

ACCESSION NUMBER: 1968:496557 HCAPLUS

DOCUMENT NUMBER: 69:96557

TITLE: New cyclic systems derived from indazole

AUTHOR(S): Portal, Carlos R.; Dennler, Enrique B.; Frasca, Adolfo R.

CORPORATE SOURCE: Univ. Buenos Aires, Buenos Aires, Argent.

SOURCE: Anales de la Asociacion Quimica Argentina (1967), 55(3-4), 245-51

CODEN: AAQAAE; ISSN: 0365-0375

DOCUMENT TYPE: Journal

LANGUAGE: Spanish

GI For diagram(s), see printed CA Issue.

AB Two new cyclic systems (I and II) were synthesized from the p-nitrophenylhydrazones of 5-acetylindane (III) by cyclization. An equimolar mixture of p-nitrophenylhydrazine and 5-acetylindane in EtOH was heated to precipitate III, m. 188-90° (EtOH). III was treated with 10-15 times its weight of polyphosphoric acid and the mixture heated to 160° 2 min., diluted with water, and extracted with C<sub>6</sub>H<sub>6</sub>, the organic phase washed with water, dried, concentrated and chromatographed on an Al<sub>2</sub>O<sub>3</sub> column using C<sub>6</sub>H<sub>6</sub> eluent. The first fraction contained 23% of a mixture of I (R = p-O<sub>2</sub>NC<sub>6</sub>H<sub>4</sub>) (IV) and II (R = p-O<sub>2</sub>NC<sub>6</sub>H<sub>4</sub>) (V). Recrystn. from EtOH yielded IV, m. 189-90°. The recrystn. liquid was evaporated and the residue chromatographed on a silica gel column with petroleum-EtOAc eluent which caused separation into 2 yellow bands. The first band was IV, and the second was V, m. 163-4° (EtOH). IV, 220 mg., was treated with 1 g. SnCl<sub>2</sub>, 4 ml. concentrate HCl, and 4 ml. AcOH and the mixture refluxed 10 min., cooled, treated with NaOH, and extracted with Et<sub>2</sub>O, and the ether phase washed with water, and dried to give I (R = p-H<sub>2</sub>NC<sub>6</sub>H<sub>4</sub>) (VI), m. 150-1° (EtOH-water). Similar treatment of V yielded II (R = p-H<sub>2</sub>NC<sub>6</sub>H<sub>4</sub>) (VII), m. 166-7° (EtOH-water). VI (240 mg.) in 11 ml. 25% H<sub>2</sub>SO<sub>4</sub> was treated with 2.4 ml. 10% Na<sub>2</sub>Cr<sub>2</sub>O<sub>7</sub> and the mixture maintained 2 hrs. at 0°, diluted with water, treated with NaOAc and extracted with Et<sub>2</sub>O and the extract washed with water, and dried to give I (R = H), m. 203-4° (Et<sub>2</sub>O-H<sub>2</sub>O). Similar treatment of V yielded II (R = H), m. 195-6° (EtOH-H<sub>2</sub>O).

IT 16640-94-1P 16641-17-1P

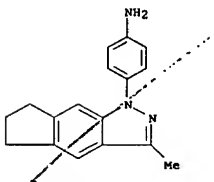
RL: SPN (Synthetic preparation); PREP (Preparation)

(preparation and spectrum (N.M.R. and uv) of)

RN 16640-94-1 HCAPLUS

CN Cyclopent[1,2]indazole, 1-(p-aminophenyl)-1,5,6,7-tetrahydro-3-methyl- (8CI)

(CA INDEX NAME)



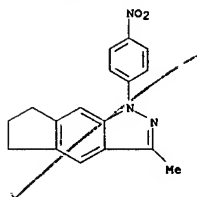
RN 16641-17-1 HCAPLUS

CN Cyclopent[1,2]indazole, 1,5,6,7-tetrahydro-3-methyl-1-(p-nitrophenyl)- (8CI)

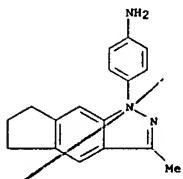
(CA INDEX NAME)

L17 ANSWER 2 OF 3 HCAPLUS COPYRIGHT 2007 ACS on STN

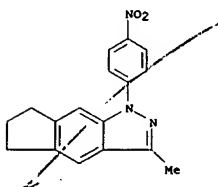
(Continued)



L17 ANSWER 3 OF 3 HCAPLUS COPYRIGHT 2007 ACS on STN  
 ACCESSION NUMBER: 1967:469259 HCAPLUS  
 DOCUMENT NUMBER: 67:69259  
 TITLE: The nuclear magnetic resonance spectra of substituted indazoles  
 AUTHOR(S): Dennier, Enrique B.; Portal, C. R.; Frasca, Adolfo R.  
 CORPORATE SOURCE: Fac. Cienc. Exact. Nat., Buenos Aires, Argent.  
 SOURCE: Spectrochimica Acta, Part A: Molecular and Biomolecular Spectroscopy (1967), 23(8), 2243-8  
 CODEN: SAMCAS; ISSN: 1386-1425  
 DOCUMENT TYPE: Journal  
 LANGUAGE: English  
 AB Proton magnetic resonance spectra of substituted indazoles, 1-p-aminophenyl-, and 1-p-nitrophenylindazoles are described and the effects of substituents on the pyrazole and benzene rings are discussed.  
 IT 16640-94-1P 16641-17-1P  
 RL: PREP (Preparation of)  
 (preparation of)  
 RN 16640-94-1 HCAPLUS  
 CN Cyclopent[f]indazole, 1-(p-aminophenyl)-1,5,6,7-tetrahydro-3-methyl- (8CI)  
 (CA INDEX NAME)



RN 16641-17-1 HCAPLUS  
 CN Cyclopent[f]indazole, 1,5,6,7-tetrahydro-3-methyl-1-(p-nitrophenyl)- (8CI)  
 (CA INDEX NAME)





Robert Havlin - 10.544899

=> log hold

COST IN U.S. DOLLARS

FULL ESTIMATED COST

DISCOUNT AMOUNTS (FOR QUALIFYING ACCOUNTS)

CA SUBSCRIBER PRICE

SINCE FILE

ENTRY

21.01

SINCE FILE

ENTRY

-2.34

TOTAL

SESSION

760.65

TOTAL

SESSION

-5.46

SESSION WILL BE HELD FOR 120 MINUTES

STN INTERNATIONAL SESSION SUSPENDED AT 10:50:08 ON 26 APR 2007

Robert Havlin - 10.544899

Connecting via Winsock to STN

Welcome to STN International! Enter x:x

LOGINID:SSPTARHH1626

PASSWORD:

\*\*\*\*\* RECONNECTED TO STN INTERNATIONAL \*\*\*\*\*  
SESSION RESUMED IN FILE 'HCAPLUS' AT 10:56:07 ON 26 APR 2007  
FILE 'HCAPLUS' ENTERED AT 10:56:07 ON 26 APR 2007  
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COST IN U.S. DOLLARS	SINCE FILE ENTRY	TOTAL SESSION
FULL ESTIMATED COST	21.01	760.65
DISCOUNT AMOUNTS (FOR QUALIFYING ACCOUNTS)	SINCE FILE ENTRY	TOTAL SESSION
CA SUBSCRIBER PRICE	-2.34	-5.46

=> file hcaplus

COST IN U.S. DOLLARS	SINCE FILE ENTRY	TOTAL SESSION
FULL ESTIMATED COST	21.01	760.65
DISCOUNT AMOUNTS (FOR QUALIFYING ACCOUNTS)	SINCE FILE ENTRY	TOTAL SESSION
CA SUBSCRIBER PRICE	-2.34	-5.46

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FILE COVERS 1907 - 26 Apr 2007 VOL 146 ISS 18  
FILE LAST UPDATED: 25 Apr 2007 (20070425/ED)

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This file contains CAS Registry Numbers for easy and accurate substance identification.

=> s us200!-544899/apps

4 US200!-544899/AP  
0 US200!-544899/PRN  
L18 4 US200!-544899/APPS  
(US200!-544899/AP,PRN)

=> d ibib

L18 ANSWER 1 OF 4 HCAPLUS COPYRIGHT 2007 ACS on STN  
ACCESSION NUMBER: 2005:1005939 HCAPLUS  
DOCUMENT NUMBER: 143:294198  
TITLE: Electrochemical fabrication process including process  
monitoring, making corrective action decisions, and  
taking appropriate actions  
INVENTOR(S): Cohen, Adam L.; Lockard, Michael S.; Smalley, Dennis  
R.; Kilgo, Marvin M.  
PATENT ASSIGNEE(S): Microfabrice Inc., USA  
SOURCE: U.S. Pat. Appl. Publ., 28 pp., Cont.-in-part of U.S.  
Ser. No. 434,494.  
CODEN: USXXCO  
DOCUMENT TYPE: Patent  
LANGUAGE: English  
FAMILY ACC. NUM. COUNT: 37  
PATENT INFORMATION:

PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
US 2005202660	A1	20050915	US 2004-995609	20041122
EP 1518822	A2	20050330	EP 2004-15983	20021206
EP 1518822	A3	20070228		
R: AT, BE, CH, DE, DK, ES, FR, GB, GR, IT, LI, LU, NL, SE, MC, PT, IE, SI, LT, LV, FI, MK, CY, AL, TR, BG, CZ, EE, SK				
US 2004000489	A1	20040101	US 2003-434494	20030507
US 2004007470	A1	20040115	US 2003-434519	20030507
AU 2003280468	A1	20040119	AU 2003-280468	20030627
US 2007045122	A1	20070301	US 2006-544899	20061006 <--
PRIORITY APPLN. INFO.:				
			US 2002-379130P	P 20020507
			US 2002-379132P	P 20020507
			US 2003-434494	A2 20030507
			US 2003-434519	A2 20030507
			US 2001-340372P	P 20011206
			US 2002-364261P	P 20020313
			US 2002-379133P	P 20020507
			US 2002-379135P	P 20020507
			US 2002-379177P	P 20020507
			US 2002-379182P	P 20020507
			US 2002-379184P	P 20020507
			US 2002-392531P	P 20020627
			US 2002-415371P	P 20021001
			US 2002-415374P	P 20021001
			US 2002-422007P	P 20021029
			US 2002-422982P	P 20021101
			US 2002-429483P	P 20021126
			US 2002-429484P	P 20021126
			US 2002-309521	A 20021203
			US 2002-430809P	P 20021203
			EP 2002-786937	A3 20021206
			US 2003-464504P	P 20030421
			US 2003-434103	A 20030507
			US 2003-434295	A 20030507
			US 2003-434497	A 20030507
			US 2003-476554P	P 20030606
			WO 2003-US20458	W 20030627

,Robert Havlin - 10.544899

=> d ibib 2

L18 ANSWER 2 OF 4 HCAPLUS COPYRIGHT 2007 ACS on STN  
 ACCESSION NUMBER: 2005:303040 HCAPLUS  
 DOCUMENT NUMBER: 142:344105  
 TITLE: Multi-step release method for electrochemically  
 fabricated structures  
 INVENTOR(S): Cohen, Adam L.; Lockard, Michael S.; McPherson, Dale  
 S.  
 PATENT ASSIGNEE(S): Microfabrics Inc., USA  
 SOURCE: U.S. Pat. Appl. Publ., 32 pp., Cont.-in-part of U.S.  
 Ser. No. 607,931.  
 CODEN: USXXCO  
 DOCUMENT TYPE: Patent  
 LANGUAGE: English  
 FAMILY ACC. NUM. COUNT: 37  
 PATENT INFORMATION:

PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
US 2005072681	A1	20050407	US 2004-841347	20040507
US 2003222738	A1	20031204	US 2002-309521	20021203
EP 1518822	A2	20050330	EP 2004-15983	20021206
EP 1518822	A3	20070228		
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US 2004004001	A1	20040108	US 2003-434295	20030507
US 2004007468	A1	20040115	US 2003-434497	20030507
US 2004007470	A1	20040115	US 2003-434519	20030507
US 2004020782	A1	20040205	US 2003-434103	20030507
US 7160429	B2	20070109		
WO 2004004061	A1	20040108	WO 2003-US20458	20030627
W: AE, AG, AL, AM, AT, AU, AZ, BA, BB, BG, BR, BY, BZ, CA, CH, CN, CO, CR, CU, CZ, DE, DK, DM, DZ, EC, EE, ES, FI, GB, GD, GE, GH, GM, HR, HU, ID, IL, IN, IS, JP, KE, KG, KP, KR, KZ, LC, LK, LR, LS, LT, LU, LV, MA, MD, MG, MK, MN, MW, MX, ME, NO, NZ, OM, PH, PL, PT, RO, RU, SC, SD, SE, SG, SK, SL, TJ, TM, TN, TR, TT, TZ, UA, UG, UZ, VC, VN, YU, ZA, ZM, ZW				
RW: GH, GM, KE, LS, MW, MZ, SD, SL, SZ, TZ, UG, ZM, ZW, AM, AZ, BY, KG, KZ, MD, RU, TJ, TM, AT, BE, BG, CH, CY, CZ, DE, DK, EE, ES, FI, FR, GB, GR, HU, IE, IT, LU, MC, NL, PT, RO, SE, SI, SK, TR, BF, BJ, CF, CG, CI, CM, GA, GN, GQ, GW, ML, MR, NE, SN, TD, TG				
AU 2003280468	A1	20040119	AU 2003-280468	20030627
CN 1669177	A	20050914	CN 2003-817237	20030627
WO 2004032210	A2	20040415	WO 2003-US31244	20031001
WO 2004032210	A3	20050609		
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RW: GH, GM, KE, LS, MW, MZ, SD, SL, SZ, TZ, UG, ZM, ZW, AM, AZ, BY, KG, KZ, MD, RU, TJ, TM, AT, BE, BG, CH, CY, CZ, DE, DK, EE, ES, FI, FR, GB, GR, HU, IE, IT, LU, MC, NL, PT, RO, SE, SI, SK, TR, BF, BJ, CF, CG, CI, CM, GA, GN, GQ, GW, ML, MR, NE, SN, TD, TG				
AU 2003279763	A1	20040423	AU 2003-279763	20031001
TW 236453	B	20050721	TW 2003-92132582	20031120
TW 244799	B	20051201	TW 2004-93116127	20040604
US 2005221529	A1	20051006	US 2005-139391	20050527
US 2007045122	A1	20070301	US 2006-544899	20061006

PRIORITY APPLN. INFO.:

US 2001-338638P P 20011203  
 US 2001-340372P P 20011206  
 US 2002-379130P P 20020507  
 US 2002-379133P P 20020507

L18 ANSWER 2 OF 4 HCAPLUS COPYRIGHT 2007 ACS on STN (Continued)  
 US 2002-379182P P 20020507  
 US 2002-379184P P 20020507  
 US 2002-392531P P 20020627  
 US 2002-415374P P 20021001  
 US 2002-309521 A2 20021203  
 US 2002-430809P P 20021203  
 US 2003-464504P P 20030421  
 US 2003-434103 A2 20030507  
 US 2003-434295 A2 20030507  
 US 2003-434497 A2 20030507  
 US 2003-434519 A2 20030507  
 US 2003-476554P P 20030606  
 US 2003-607931 A2 20030627  
 US 2002-364261P P 20020313  
 US 2002-379135P P 20020507  
 US 2002-379177P P 20020507  
 US 2002-415371P P 20021001  
 US 2002-422007P P 20021029  
 US 2002-422982P P 20021101  
 US 2002-429483P P 20021126  
 US 2002-429484P P 20021126  
 EP 2002-786937 A3 20021206  
 US 2002-313795 A1 20021206  
 WO 2003-US20458 W 20030627  
 WO 2003-US31244 W 20031001

,Robert Havlin - 10.544899

=> d ibib 3

L18 ANSWER 3 OF 4 HCAPLUS COPYRIGHT 2007 ACS on STN  
ACCESSION NUMBER: 2005:98637 HCAPLUS  
DOCUMENT NUMBER: 142:187861  
TITLE: Methods for electrochemically fabricating multi-layer  
structures including regions incorporating maskless,  
patterned, multiple layer thickness depositions of  
selected materials  
INVENTOR(S): Cohen, Adam L.; Smalley, Dennis R.  
PATENT ASSIGNEE(S): Microfabrica Inc., USA  
SOURCE: U.S. Pat. Appl. Publ., 35 pp., Cont.-in-part of U.S.  
Ser. No. 434,519.  
CODEN: USXXCO  
DOCUMENT TYPE: Patent  
LANGUAGE: English  
FAMILY ACC. NUM. COUNT: 37  
PATENT INFORMATION:

PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
US 2005023144	A1	20050203	US 2004-841001	20040507
EP 1518822	A2	20050330	EP 2004-15983	20021206
EP 1518822	A3	20070228		
R: AT, BE, CH, DE, DK, ES, FR, GB, GR, IT, LI, LU, NL, SE, MC, PT, IE, SI, LT, LV, FI, MK, CY, AL, TR, BG, CZ, EE, SK				
US 2004007470	A1	20040115	US 2003-434519	20030507
AU 2003280468	A1	20040119	AU 2003-280468	20030627
US 2007045122	A1	20070301	US 2006-544899	20061006 <--
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			US 2002-379130P	P 20020507
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			US 2002-415371P	P 20021001
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			US 2002-422007P	P 20021029
			US 2002-422982P	P 20021101
			US 2002-429483P	P 20021126
			US 2002-429484P	P 20021126
			US 2002-309521	A 20021203
			US 2002-430809P	P 20021203
			EP 2002-786937	A3 20021206
			US 2003-464504P	P 20030421
			US 2003-434103	A 20030507
			US 2003-434295	A 20030507
			US 2003-434497	A 20030507
			US 2003-476554P	P 20030606
			WO 2003-US20458	W 20030627

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Robert Havlin - 10.544899

=> d ibib 4





.Robert Havlin - 10.544899

=> file reg

COST IN U.S. DOLLARS	SINCE FILE ENTRY	TOTAL SESSION
FULL ESTIMATED COST	9.92	770.57
DISCOUNT AMOUNTS (FOR QUALIFYING ACCOUNTS)	SINCE FILE ENTRY	TOTAL SESSION
CA SUBSCRIBER PRICE	0.00	-5.46

FILE 'REGISTRY' ENTERED AT 10:57:13 ON 26 APR 2007  
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STRUCTURE FILE UPDATES: 25 APR 2007 HIGHEST RN 932710-95-7  
DICTIONARY FILE UPDATES: 25 APR 2007 HIGHEST RN 932710-95-7

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TSCA INFORMATION NOW CURRENT THROUGH December 2, 2006

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predicted properties as well as tags indicating availability of  
experimental property data in the original document. For information  
on property searching in REGISTRY, refer to:

<http://www.cas.org/support/stngen/stndoc/properties.html>

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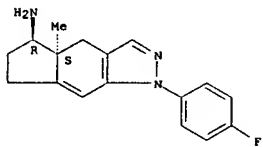
L19 TRANSFER L18 4 RN : 282 TERMS  
L20 282 L19

=> d scan

Robert Havlin - 10.544899

L20 282 ANSWERS REGISTRY COPYRIGHT 2007 ACS on STN  
IN Cyclopent[*f*]indazol-5-amine, 1-(4-fluorophenyl)-1,4,4a,5,6,7-hexahydro-4a-  
methyl-, (4a*S*,5*R*)- (9CI)  
MF C17 H18 F N3

Absolute stereochemistry.



\*\*PROPERTY DATA AVAILABLE IN THE 'PROP' FORMAT\*\*

HOW MANY MORE ANSWERS DO YOU WISH TO SCAN? (1):0

=> file hcaplus

COST IN U.S. DOLLARS	SINCE FILE ENTRY	TOTAL SESSION
FULL ESTIMATED COST	0.45	785.27
DISCOUNT AMOUNTS (FOR QUALIFYING ACCOUNTS)		
CA SUBSCRIBER PRICE	0.00	-5.46

FILE 'HCAPLUS' ENTERED AT 10:57:38 ON 26 APR 2007  
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FILE COVERS 1907 - 26 Apr 2007 VOL 146 ISS 18  
FILE LAST UPDATED: 25 Apr 2007 (20070425/ED)

New CAS Information Use Policies, enter HELP USAGETERMS for details.

This file contains CAS Registry Numbers for easy and accurate substance identification.

=> s l20  
L21 408632 L20

=> file reg

COST IN U.S. DOLLARS	SINCE FILE ENTRY	TOTAL SESSION
FULL ESTIMATED COST	2.60	787.87
DISCOUNT AMOUNTS (FOR QUALIFYING ACCOUNTS)		
CA SUBSCRIBER PRICE	0.00	-5.46

FILE 'REGISTRY' ENTERED AT 10:57:50 ON 26 APR 2007  
USE IS SUBJECT TO THE TERMS OF YOUR STN CUSTOMER AGREEMENT.  
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STRUCTURE FILE UPDATES: 25 APR 2007 HIGHEST RN 932710-95-7  
DICTIONARY FILE UPDATES: 25 APR 2007 HIGHEST RN 932710-95-7

New CAS Information Use Policies, enter HELP USAGETERMS for details.

TSCA INFORMATION NOW CURRENT THROUGH December 2, 2006

Please note that search-term pricing does apply when conducting SmartSELECT searches.

REGISTRY includes numerically searchable data for experimental and predicted properties as well as tags indicating availability of experimental property data in the original document. For information on property searching in REGISTRY, refer to:

<http://www.cas.org/support/stngen/stndoc/properties.html>

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20800526 C>=16

18452292 N>=2

3718411 F>=1

L22 262 L20 AND C>=16 AND N>=2 AND F>=1

=> file hcaplus

COST IN U.S. DOLLARS

SINCE FILE

TOTAL

ENTRY

SESSION

FULL ESTIMATED COST

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803.62

DISCOUNT AMOUNTS (FOR QUALIFYING ACCOUNTS)

SINCE FILE

TOTAL

ENTRY

SESSION

CA SUBSCRIBER PRICE

0.00

-5.46

FILE 'HCAPLUS' ENTERED AT 10:58:52 ON 26 APR 2007

USE IS SUBJECT TO THE TERMS OF YOUR STN CUSTOMER AGREEMENT.

PLEASE SEE "HELP USAGETERMS" FOR DETAILS.

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FILE COVERS 1907 - 26 Apr 2007 VOL 146 ISS 18

FILE LAST UPDATED: 25 Apr 2007 (20070425/ED)

New CAS Information Use Policies, enter HELP USAGETERMS for details.

This file contains CAS Registry Numbers for easy and accurate substance identification.

=> s l22

L23 7 L22

=> d hist

(FILE 'HOME' ENTERED AT 10:26:58 ON 26 APR 2007)

FILE 'REGISTRY' ENTERED AT 10:27:10 ON 26 APR 2007

L1 STRUCTURE UPLOADED

L2 14 S L1 SSS SAM

FILE 'HCAPLUS' ENTERED AT 10:28:33 ON 26 APR 2007

L3 3 S L2

FILE 'REGISTRY' ENTERED AT 10:32:56 ON 26 APR 2007

L4 272 S L1 SSS FULL

FILE 'HCAPLUS' ENTERED AT 10:33:15 ON 26 APR 2007

L5 4 S L4

L6 1 S L5 NOT L3

FILE 'REGISTRY' ENTERED AT 10:45:22 ON 26 APR 2007

L7 STRUCTURE UPLOADED

L8 194 S L7 SSS FULL

FILE 'HCAPLUS' ENTERED AT 10:46:02 ON 26 APR 2007

L9 4 S L8

L10 0 S L9 NOT L5

FILE 'REGISTRY' ENTERED AT 10:47:24 ON 26 APR 2007

L11 STRUCTURE UPLOADED

L12 273 S L11 SSS FULL

L13 STRUCTURE UPLOADED

L14 14 S L13 SSS SAM

L15 277 S L13 SSS FULL

FILE 'HCAPLUS' ENTERED AT 10:49:13 ON 26 APR 2007

L16 7 S L15

L17 3 S L16 NOT L5

FILE 'HCAPLUS' ENTERED AT 10:56:14 ON 26 APR 2007

L18 4 S US200!-544899/APPS

FILE 'REGISTRY' ENTERED AT 10:57:13 ON 26 APR 2007

FILE 'HCAPLUS' ENTERED AT 10:57:22 ON 26 APR 2007

L19 TRA L18 4 RN : 282 TERMS

FILE 'REGISTRY' ENTERED AT 10:57:23 ON 26 APR 2007

L20 282 SEA L19

FILE 'HCAPLUS' ENTERED AT 10:57:38 ON 26 APR 2007

L21 408632 S L20

FILE 'REGISTRY' ENTERED AT 10:57:50 ON 26 APR 2007

L22 262 S L20 AND C>=16 AND N>=2 AND F>=1

FILE 'HCAPLUS' ENTERED AT 10:58:52 ON 26 APR 2007

L23 7 S L22

=> s l23 not l16

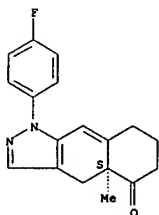
L24 3 L23 NOT L16

=> d ibib abs hitstr tot

L24 ANSWER 1 OF 3 HCAPLUS COPYRIGHT 2007 ACS on STN

ACCESSION NUMBER: 2004:767304 HCAPLUS  
 DOCUMENT NUMBER: 141:405644  
 TITLE: Design and evaluation of novel nonsteroidal dissociating glucocorticoid receptor ligands  
 AUTHOR(S): Shah, Nilesh; Scanlan, Thomas S.  
 CORPORATE SOURCE: Graduate Group in Biophysics, University of California, San Francisco, CA, 94143-2280, USA  
 SOURCE: Bioorganic & Medicinal Chemistry Letters (2004), 14(20), 5199-5203  
 CODEN: BMCLE8; ISSN: 0960-894X  
 PUBLISHER: Elsevier B.V.  
 DOCUMENT TYPE: Journal  
 LANGUAGE: English  
 OTHER SOURCE(S): CASREACT 141:405644  
 AB A novel class of phenylpyrazole fused Wieland-Miescher ketone deriva. are high affinity, receptor specific, selective modulators of glucocorticoid receptor (GR) mediated transcription in vitro, dissociating transactivation, AP-1 repression, and NF- $\kappa$ B repression from each other.  
 IT 614763-00-7P  
 RL: PRP (Properties); RCT (Reactant); SPN (Synthetic preparation); PREP (Preparation); RACT (Reactant or reagent)  
 (design and evaluation of novel nonsteroidal dissociating glucocorticoid receptor ligands)  
 RN 614763-00-7 HCAPLUS  
 CN 5H-Benz[*f*]indazol-5-one, 1-(4-fluorophenyl)-1,4,4a,6,7,8-hexahydro-4a-methyl-, (4aS)- (9CI) (CA INDEX NAME)

Absolute stereochemistry.

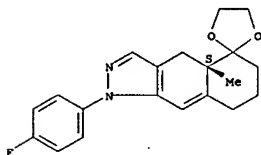


REFERENCE COUNT: 32 THERE ARE 32 CITED REFERENCES AVAILABLE FOR THIS RECORD. ALL CITATIONS AVAILABLE IN THE RE FORMAT

L24 ANSWER 2 OF 3 HCAPLUS COPYRIGHT 2007 ACS on STN (Continued)

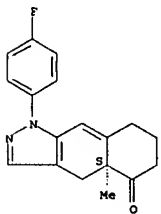
AB Octahydro-2-H-naphtho[1,2-*f*]indole-4-carboxamide deriva. I (X = CO, NHCO, CONH, NH, CH<sub>2</sub>NH; R<sub>1</sub>, R<sub>2</sub> = H, alkyl, alkenyl, cycloalkyl, alkoxy, aryl; R<sub>3</sub> = alkyl, alkoxy, acid, halogen substituted alkyl; R<sub>4</sub> = alkyl, alkenyl, cycloalkoxy, alkoxy, aryl) were prepared as selective glucocorticoid receptor modulators for the treatment of autoimmune and inflammatory conditions. Thus, (S)-Wieland-Miescher ketone was protected as the ketal using p-toluene sulfonic acid and ethylene glycol and then treated with Et formate to give the hydroxymethylene ketal derivative. The hydroxymethylene was dissolved in acetic acid and reacted with p-fluorophenyl hydrazine hydrochloride to give II. The ketal of II was converted to the ketone using 6N HCl, and the resulting ketone transformed into the triflate. The triflate was treated with tributylvinyl tin and PPh<sub>3</sub> to give the corresponding coupling product. Treatment with ethyl-4,4,4-trifluorocrotonate followed by dropwise addition of BCl<sub>3</sub> gave the target I (R<sub>1</sub> = CF<sub>3</sub>, R<sub>2</sub>, R<sub>3</sub> = H, X = CO, R<sub>4</sub> = OEt).  
 IT 614762-99-1P 614763-00-7P  
 RL: RCT (Reactant); SPN (Synthetic preparation); PREP (Preparation); RACT (Reactant or reagent)  
 (preparation of octahydronaphthoindole-4-carboxamide deriva. as selective glucocorticoid receptor modulators for the treatment of autoimmune and inflammatory conditions)  
 RN 614762-99-1 HCAPLUS  
 CN Spiro[5H-benz[*f*]indazole-5,2'-[1,3]dioxolane], 1-(4-fluorophenyl)-1,4,4a,6,7,8-hexahydro-4a-methyl-, (4aS)- (9CI) (CA INDEX NAME)

Absolute stereochemistry.



RN 614763-00-7 HCAPLUS  
 CN 5H-Benz[*f*]indazol-5-one, 1-(4-fluorophenyl)-1,4,4a,6,7,8-hexahydro-4a-methyl-, (4aS)- (9CI) (CA INDEX NAME)

Absolute stereochemistry.

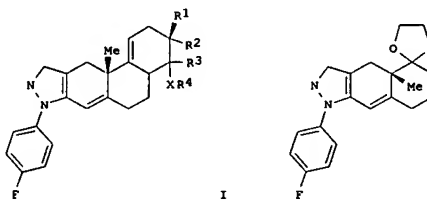


L24 ANSWER 2 OF 3 HCAPLUS COPYRIGHT 2007 ACS on STN

ACCESSION NUMBER: 2004:270010 HCAPLUS  
 DOCUMENT NUMBER: 140:287380  
 TITLE: Preparation of octahydro-2-H-naphtho[1,2-*f*]indole-4-carboxamide derivatives as selective glucocorticoid receptor modulators for the treatment of autoimmune and inflammatory conditions  
 INVENTOR(S): Ali, Amjad; Aster, Susan D.; Balkovec, James M.; Graham, Donald W.; Hunt, Julianne A.; Kallashi, Florida; Sinclair, Peter J.; Tata, James R.; Taylor, Gayle E.; Goulet, Joung L.  
 PATENT ASSIGNEE(S): Merck & Co., Inc., USA  
 SOURCE: PCT Int. Appl., 170 pp.  
 CODEN: PIXXD2  
 DOCUMENT TYPE: Patent  
 LANGUAGE: English  
 FAMILY ACC. NUM. COUNT: 1  
 PATENT INFORMATION:

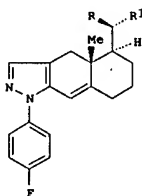
PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
WO 2004026248	A2	20040401	WO 2003-US29494	20030917
WO 2004026248	A3	20040715		
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RW:	GH, GM, KE, LS, MW, MZ, SD, SL, SZ, TZ, UG, ZM, ZW, AM, AZ, BY, KG, KZ, MD, RU, TJ, TM, AT, BE, BG, CH, CY, CZ, DE, DK, EE, ES, FI, FR, GB, GR, HU, IE, IT, LU, MC, NL, PT, RO, SE, SI, SK, TR, BF, BJ, CF, CG, CI, CM, GA, GN, GQ, GW, ML, MR, NE, SN, TD, TG			
CA 2499150	A1	20040401	CA 2003-2499150	20030917
AU 2003270783	A1	20040409	AU 2003-270783	20030917
EP 1542996	A2	20050622	EP 2003-752495	20030917
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JP 2006503107	T	20060126	JP 2004-568945	20030917
US 2005245588	A1	20051103	US 2005-527615	20050311
PRIORITY APPLN. INFO.:			US 2002-412231P	P 20020920
			US 2003-476079P	P 20030605
			WO 2003-US29494	W 20030917

OTHER SOURCE(S): MARPAT 140:287380  
 GI



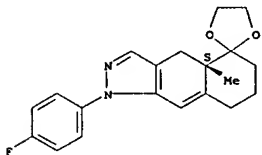
L24 ANSWER 3 OF 3 HCAPLUS COPYRIGHT 2007 ACS on STN

ACCESSION NUMBER: 2004:267662 HCAPLUS  
 DOCUMENT NUMBER: 141:7063  
 TITLE: Novel N-Arylpyrazolo[3,2-*c*]-Based Ligands for the Glucocorticoid Receptor: Receptor Binding and in Vivo Activity  
 AUTHOR(S): Ali, Amjad; Thompson, Christopher F.; Balkovec, James M.; Graham, Donald W.; Hammond, Milton L.; Quraishi, Nazia; Tata, James R.; Einstein, Monica; Ge, Len; Harris, Georgianna; Kelly, Terri M.; Mazur, Paul; Pandit, Shipra; Santoro, Joseph; Sitlani, Ayesha; Wang, Chuanlin; Williamson, Joanne; Miller, Douglas K.; Thompson, Chris M.; Zaller, Dennis M.; Forrest, Michael J.; Carballea-Jane, Ester; Luell, Silvi  
 CORPORATE SOURCE: Departments of Medicinal Chemistry, Metabolic Disorders Immunology and Pharmacology, Merck Research Laboratories, Rahway, NJ, 07065, USA  
 SOURCE: Journal of Medicinal Chemistry (2004), 47(10), 2441-2452  
 CODEN: JMCMAR; ISSN: 0022-2623  
 PUBLISHER: American Chemical Society  
 DOCUMENT TYPE: Journal  
 LANGUAGE: English  
 OTHER SOURCE(S): CASREACT 141:7063  
 GI



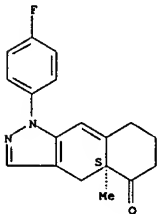
AB A novel series of selective ligands for the human glucocorticoid receptor (hGR) are described. Preliminary structure-activity relationships were focused on substitution at C-1 and indicated a preference for 3-, 4-, and 5-substituted aromatic and benzyl groups. The resulting analogs, e.g., I [R = OH, R<sub>1</sub> = 3,4,5-MeO(F)<sub>2</sub>CGH<sub>2</sub>, CH<sub>2</sub>CGH<sub>2</sub>FP-4], exhibited excellent affinity for hGR (IC<sub>50</sub> 1.9 nM and 2.8 nM, resp.) and an interesting partial agonist profile in functional assays of transactivation (tyrosine aminotransferase, TAT, and glutamine synthetase, GS) and transrepression (IL-6). The most potent compds. were I [R = 4-FC6H<sub>4</sub>, 2-thienyl, R<sub>1</sub> = OH]. These candidates showed highly efficacious IL-6 inhibition vs. dexamethasone. I [R = 2-thienyl, R<sub>1</sub> = OH] was evaluated in vivo in the mouse LPS challenge model and showed an ED<sub>50</sub> = 4.0 mg/kg, compared to 0.5 mg/kg for prednisolone in the same assay.  
 IT 614762-99-1P 614763-00-7P  
 RL: RCT (Reactant); SPN (Synthetic preparation); PREP (Preparation); RACT (Reactant or reagent)  
 (preparation and glucocorticoid receptor binding of [aryl(hydroxy)methyl]naphthopyrazoles)  
 RN 614762-99-1 HCAPLUS  
 CN Spiro[5H-benz[*f*]indazole-5,2'-[1,3]dioxolane], 1-(4-fluorophenyl)-1,4,4a,6,7,8-hexahydro-4a-methyl-, (4aS)- (9CI) (CA INDEX NAME)

Absolute stereochemistry.



RN 614763-00-7 HCAPLUS  
CN 5H-Benz[f]indazol-5-one, 1-(4-fluorophenyl)-1,4,4a,6,7,8-hexahydro-4a-methyl-, (4aS)- (9CI) (CA INDEX NAME)

Absolute stereochemistry.



REFERENCE COUNT: 36 THERE ARE 36 CITED REFERENCES AVAILABLE FOR THIS RECORD. ALL CITATIONS AVAILABLE IN THE RE FORMAT